

GenCore version 4.5  
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OM protein - protein search, using sw model

Run on: April 25, 2000, 20:10:14 ; Search time 85.2 Seconds  
(without alignments)  
176.812 Million cell updates/sec

Title: US-09-125-005-6

Perfect score: 3384

Sequence: 1 MAQSTATSPDGGTTFEHLWS.....PDCKARKQPIKEEFAEIH 636

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 188963 seqs, 23686106 residues

Total number of hits satisfying chosen parameters: 188963

Minimum DB seq length: 0

Maximum DB seq length: 1000000

Post-processing: Minimum Match 0%

Listing first 45 summaries

Database : A\_Geneseq\_36.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	3384	100.0	636	1 W36184	Human p53 tumour s
2	3367.5	99.5	635	1 W30661	Human NBS-1 alpha
3	3304.5	97.7	637	1 W36182	Monkey p53 tumour
4	3330	92.5	588	1 W36189	Human p53 tumour s
5	3058	90.4	587	1 W36187	Human p53 tumour s
6	2830.5	83.6	589	1 W36185	Mouse p53 tumour s
7	2624	77.5	499	1 W36190	Human p53 tumour s
8	2574	76.1	499	1 W36183	Monkey p53 tumour
9	2330.5	68.9	506	1 W36188	Human p53 tumour s
10	726.5	21.5	401	1 W28487	Human p53 protein
11	725.5	21.4	401	1 W28488	Human p53 protein
12	724.5	21.4	355	1 W13950	Del356-393 modifie
13	724.5	21.4	363	1 W13954	Modified p53 varia
14	724.5	21.4	393	1 R22238	Sequence of 53 KD
15	724.5	21.4	393	1 R26738	p53. Synthetic onc
16	724.5	21.4	393	1 R79658	Human p53 protein.
17	724.5	21.4	393	1 R94623	p53 protein. Recom
18	724.5	21.4	393	1 R91933	Wild type p53 prot
19	724.5	21.4	393	1 W02617	Human p53 tumour s
20	724.5	21.4	393	1 W05344	Human p53. New hum
21	724.5	21.4	393	1 W05348	Human wild-type p5
22	724.5	21.4	393	1 W57242	Human p53 protein
23	724.5	21.4	393	1 W57243	Human p53 protein
24	724.5	21.4	393	1 W48658	Amino acid sequenc
25	724.5	21.4	393	1 W69217	Human wild-type p5
26	724.5	21.4	393	1 W69718	Human p53 used in
27	724.5	21.4	393	1 Y03191	Amino acid sequenc
28	724.5	21.4	438	1 R74272	Tumour suppressor
29	724.5	21.4	533	1 W19763	p53-GM-CSF immuno
30	723.5	21.4	363	1 W13971	Modified p53 varia
31	723.5	21.4	393	1 W13949	T284R modified hum
32	723.5	21.4	393	1 W13953	T284K modified hum
33	723.5	21.4	393	1 W57244	Human p53 protein
34	723.5	21.4	393	1 W57245	Human p53 protein

35 723.5 21.4 393 1 W84270 Human p53 protein.  
36 723.5 21.4 438 1 R50088 p53 tumour suppress  
37 720.5 21.3 363 1 W13972 Modified p53 varia  
38 720.5 21.3 393 1 W05347 Human p53 mutant R  
39 720.5 21.3 393 1 W13951 Human tumour-deriv  
40 720 21.3 354 1 R51874 Human p53 amino ac  
41 719.5 21.3 363 1 W13973 Modified p53 varia  
42 719.5 21.3 363 1 W13974 Modified p53 varia  
43 719.5 21.3 393 1 W05345 Human p53 mutant N  
44 719.5 21.3 393 1 W05346 Human p53 mutant R  
45 719.5 21.3 393 1 W13968 Modified p53 varia

#### ALIGNMENTS

RESULT 1

ID W36184 standard; Protein; 636 AA.  
AC W36184; 1998 (first entry)  
DE Human p53 tumour suppressor-related protein SR-p70a.  
KW SR-p70; human; transcription factor; p53; tumour suppressor gene;  
KW homology; differential splicing; diagnosis; cancer; neuroblastoma;  
KW gene therapy; apoptosis.  
OS Homo sapiens.  
PN W09728186-A1.  
PD 07-AUG-1997.  
PF 02-FEB-1997; F00214.  
PR 02-FEB-1996; FR-001309.  
PA (SNFI ) SANOFI SA.  
PI Caput D, Fexarava P, Kaghad AM;  
DR WPI; 97-402550/37.  
DR N-PSDB; V01498.  
PT New polypeptide(s) encoded by the SR-p70 tumour suppressor gene -  
PT and related nucleic acid, useful for diagnosis and treatment of  
PT tumours  
PS Claim 7: Fig 6: 136pp; French.  
CC This is the amino acid sequence of the human protein SR-p70a. SR-p70  
CC are transcription factors which may control the activity of p53-regulated  
CC genes, and are expressed by tumour suppressor genes related to the p53  
CC gene family. The gene sequence was isolated from the human colon  
CC adenocarcinoma cell line Ht-29, using primers V01506-7. The sequence  
CC can be used in the diagnosis and monitoring of cancer, especially  
CC neuroblastoma. The nucleic acid sequences and corresponding antisense  
CC sequences, are also useful in gene therapy, e.g. to regulate apoptosis.  
SQ Sequence 636 AA;

Query Match 100.0%; Score 3384; DB 1; Length 636;  
Best Local Similarity 100.0%; Pred. No. 1.1e-287; Mismatches 0; Indels 0; Gaps 0;  
Matches 636; Conservative 0;  
QY 1 MAQSTATSPDGGTTFEHLWSLEPDSSTYFDLPQSSRGNNVEVGGTSSMDVFLHCGMTTS 60  
Db 1 MAQSTATSPDGGTTFEHLWSLEPDSSTYFDLPQSSRGNNVEVGGTSSMDVFLHCGMTTS 60  
QY 61 VMAQFNLLSTMDQMSRAASAPYTPPEHAASVTPHSYPAQPSSTFTDTSAPVIPSNTD 120  
Db 61 VMAQFNLLSTMDQMSRAASAPYTPPEHAASVTPHSYPAQPSSTFTDTSAPVIPSNTD 120  
QY 121 YPGPHPEVTFQSSSTAKSATWTYSPLLKLYCOIAKTCPIQIKVSTPPPGTAIRAMPV 180  
Db 121 YPGPHPEVTFQSSSTAKSATWTYSPLLKLYCOIAKTCPIQIKVSTPPPGTAIRAMPV 180  
QY 181 YKKAHVTDVVKPCPNHELGRDENEGQSAPASHLIRVEGNLSQYVDDPVTGRQSVVVPY 240  
Db 181 YKKAHVTDVVKPCPNHELGRDENEGQSAPASHLIRVEGNLSQYVDDPVTGRQSVVVPY 240  
QY 241 EPPQVGTEFTTILYNFMCNCSVCVGGMNRRLPILITLMDRDQVGLGRSFEGRICACPR 300  
Db 241 EPPQVGTEFTTILYNFMCNCSVCVGGMNRRLPILITLMDRDQVGLGRSFEGRICACPR 300



Query Match 97.7%; Score 3304.5; DB 1; Length 637;  
 Best Local Similarity 97.5%; Pred. No. 1e-280;  
 Matches 621; Conservative 4; Mismatches 11; Indels 1; Gaps 1;

QY 1 MAQSTATSPDGGTTFFHLWSSLEPDSYFDLPQSSRGNEVYGGTSSMDVFLHSGMTS 60  
 DB 1 MAQSTTSPDGGTTFFHLWSSLEPDSYFDLPQSSRGNEVYGGTSSMDVFLHSGMTS 60

QY 61 VMAQFLLSTMDQSSRAASAPYTPHEAASVTPHSPYAQSSFTDTMSPAPVPSNTD 120  
 DB 61 VMAQFLLSTMDQSSRAASAPYTPHEAASVTPHSPYAQSSFTDTMSPAPVPSNTD 120

QY 121 YPGPHFEVTFQSSSTAKSATWTYSPLLLKLYCQIAKTCPIQIKVSTPPPGTAIRAMPV 180  
 DB 121 YPGPHFEVTFQSSSTAKSATWTYSPLLLKLYCQIAKTCPIQIKVSTPPPGTAIRAMPV 180

QY 181 YKAEHVTDVYKRCNHELGRDFNEGOSAPASHLIRVEGNLSQYVDDPVTGRQSVVY 240  
 DB 181 YKAEHVTDVYKRCNHELGRDFNEGOSAPASHLIRVEGNLSQYVDDPVTGRQSVVY 240

QY 241 EPPQVGTETTYLYNFMCHSCVGGMNRRLPIIIITLEMRDQGVLRGRSFEGRICACPG 300  
 DB 241 EPPQVGTETTYLYNFMCHSCVGGMNRRLPIIIITLEMRDQGVLRGRSFEGRICACPG 300

QY 301 DRKADEHYREOQALNESSAKNGAASKRAFQKSPAPVAPALGAGYKRRRGDEDTYLYQVR 360  
 DB 301 DRKADEHYREOQALNESSAKNGAASKRAFQKSPAPVAPALGAGYKRRRGDEDTYLYQVR 360

QY 361 GRENFIILMKLESLELMELVPOPLVDSYRQOQLLQRPSSHLPQPSYGPVLSPMKNVHG 420  
 DB 361 GRENFIILMKLESLELMELVPOPLVDSYRQOQLLQRPSSHLPQPSYGPVLSPMKNVHG 420

QY 421 MNKLPVNLVCGOPPHSSAATPNLGPVGMNNHGHAVPANGEMSSSHSAQSVSGSH 480  
 DB 421 MNKLPVNLVCGOPPHSSAATPNLGPVGMNNHGHAVPANGEMSSSHSAQSVSGSH 480

QY 481 CTTPPPYHADPSLVSLTGLGCPNCIEYFTSQGLSIYHLQNTIEDLGALKIPEQYRMT 540  
 DB 481 CTTPPPYHADPSLVSLTGLGCPNCIEYFTSQGLSIYHLQNTIEDLGALKIPEQYRMT 540

QY 541 IWRGLQDLKQGHY-STAQQLRSSNAATISIGSGELQQRVMEAVHFRVHTTIPNR 599  
 DB 541 IWRGLQDLKQGHY-STAQQLRSSNAATISIGSGELQQRVMEAVHFRVHTTIPNR 600

QY 600 GPGGGPDWADFGDLPCKARKQPIKEEFTAEI 636  
 DB 601 GPGGAGDEWADFGDLPCKARKQPIKEEFTAEI 637

RESULT 4  
 W36189  
 ID W36189 standard; Protein; 588 AA.  
 AC W36189;  
 DT 27-APR-1998 (first entry)  
 DE Human p53 tumour suppressor-related protein SR-p70f.  
 KW SR-p70; human; transcription factor; p53; tumour suppressor gene;  
 KW homology; differential splicing; diagnosis; cancer; neuroblastoma;  
 KW gene therapy; apoptosis.  
 OS Homo sapiens.  
 PN W09728186-AL.  
 PD 07-AUG-1997.  
 PF 03-FEB-1997; F00214.  
 PR 02-FEB-1996; FR-001309.  
 PA (SNEI) SANOFI SA.  
 PI Caput D, Ferrara P, Kaghad AM;  
 DR WPI; 97-402550/37.  
 DR N-PSDB; V01504.  
 PT New polypeptide(s) encoded by the SR-p70 tumour suppressor gene  
 PT and related nucleic acid, useful for diagnosis and treatment of  
 PT tumours  
 PS Claim 1; Page 69-70; 135pp; French.  
 CC This is the amino acid sequence of the human protein SR-p70f. SR-p70 are  
 CC transcription factors which may control the activity of p53-regulated

CC genes, and are expressed by tumour suppressor genes related to the p53  
 CC gene family. The gene sequence was isolated from the human neuroblastoma  
 CC cell line SR-N-SH, using primers V01515 and V01518. The SR-p70f gene  
 CC sequence contains a 98 bp deletion between bases 24-25 as compared to  
 CC the SR-p70a sequence (V01498). This deletion causes a loss of the  
 CC translation initiation codon found in SR-p70a, resulting in the use of  
 CC a downstream AUG (corresponding to an internal Met codon in SR-p70a).  
 CC The protein is truncated by 48 amino acids at the N-terminus as compared  
 CC to the SR-p70a protein (W36184). The sequence can be used in the  
 CC diagnosis and monitoring of cancer, especially neuroblastoma. The  
 CC nucleic acid sequences and corresponding antisense sequences, are also  
 CC useful in gene therapy, e.g. to regulate apoptosis.  
 SQ Sequence 588 AA;

Query Match 92.5%; Score 3130; DB 1; Length 588;  
 Best Local Similarity 100.0%; Pred. No. 1.7e-265;  
 Matches 588; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 49 MDVFLHSGMTSVMAQFLLSTMDQSSRAASAPYTPHEAASVTPHSPYAQSSFTDT 108  
 DB 1 MDVFLHSGMTSVMAQFLLSTMDQSSRAASAPYTPHEAASVTPHSPYAQSSFTDT 60

QY 109 MSPAPVPSNTDYPGPHFEVTFQSSSTAKSATWTYSPLLLKLYCQIAKTCPIQIKVSTP 168  
 DB 61 MSPAPVPSNTDYPGPHFEVTFQSSSTAKSATWTYSPLLLKLYCQIAKTCPIQIKVSTP 120

QY 169 PPGTATRAMPVYKAEHVTDVYKRCNHELGRDFNEGOSAPASHLIRVEGNLSQYVDD 228  
 DB 121 PPGTATRAMPVYKAEHVTDVYKRCNHELGRDFNEGOSAPASHLIRVEGNLSQYVDD 180

QY 229 PVTGRQSVVYVPGVGTETTYLYNFMCHSCVGGMNRRLPIIIITLEMRDQGVLRGR 288  
 DB 181 PVTGRQSVVYVPGVGTETTYLYNFMCHSCVGGMNRRLPIIIITLEMRDQGVLRGR 240

QY 289 SFEGRICACGRDKADEHYREOQALNESSAKNGAASKRAFQKSPAPVAPALGAGYKRRR 348  
 DB 241 SFEGRICACGRDKADEHYREOQALNESSAKNGAASKRAFQKSPAPVAPALGAGYKRRR 300

QY 349 HGDEDTYLYQVRGRENFEILMKLESLELMELVPOPLVDSYRQOQLLQRPSSHLPQPSY 408  
 DB 301 HGDEDTYLYQVRGRENFEILMKLESLELMELVPOPLVDSYRQOQLLQRPSSHLPQPSY 360

QY 409 PVLSPMKNVHGGMKNLPSVNLVCGOPPHSSAATPNLGPVGMNNHGHAVPANGEMSS 468  
 DB 361 PVLSPMKNVHGGMKNLPSVNLVCGOPPHSSAATPNLGPVGMNNHGHAVPANGEMSS 420

QY 469 SHSAQSVSGSHCTPPPPYHADPSLVSLTGLGCPNCIEYFTSQGLSIYHLQNTIEDL 528  
 DB 421 SHSAQSVSGSHCTPPPPYHADPSLVSLTGLGCPNCIEYFTSQGLSIYHLQNTIEDL 480

QY 529 GALKIPEQYRMTIWRGLQDLKQGHY-STAQQLRSSNAATISIGSGELQQRVMEAVHFR 588  
 DB 481 GALKIPEQYRMTIWRGLQDLKQGHY-STAQQLRSSNAATISIGSGELQQRVMEAVHFR 540

QY 589 RVHRTTIPNRGGPGGPDWADFGDLPCKARKQPIKEEFTAEI 636  
 DB 541 RVHRTTIPNRGGPGGPDWADFGDLPCKARKQPIKEEFTAEI 588

RESULT 5  
 W36187  
 ID W36187 standard; Protein; 587 AA.  
 AC W36187;  
 DT 27-APR-1998 (first entry)  
 DE Human p53 tumour suppressor-related protein SR-p70d.  
 KW SR-p70; human; transcription factor; p53; tumour suppressor gene;  
 KW homology; differential splicing; diagnosis; cancer; neuroblastoma;  
 KW gene therapy; apoptosis.  
 OS Homo sapiens.  
 PN W09728186-AL.  
 PD 07-AUG-1997.  
 PF 03-FEB-1997; F00214.

DE Mouse p53 tumour suppressor-related protein SR-p70c.  
 KW SR-p70; mouse; transcription factor; p53; tumour suppressor gene;  
 KW homology; differential splicing; diagnosis; cancer; neuroblastoma;  
 KW gene therapy; apoptosis.  
 PS Mus musculus.  
 PN W09728186-A1.  
 PD 07-AUG-1997.  
 PF 03-FEB-1997; F00214.  
 PR 02-FEB-1996; FR-001309.  
 PA (SNFI) SANOFI SA.  
 PI Caput D, Ferrara P, Kaghad AM;  
 DR WPI; 97-402550/37.  
 DR N-PSDB; V01502.  
 PT New polypeptide(s) encoded by the SR-p70 tumour suppressor gene -  
 and related nucleic acid, useful for diagnosis and treatment of  
 tumours  
 PS Claim 1; Page 62-64; 136pp; French.  
 CC This is the amino acid sequence of the human protein SR-p70d. R-p70 are  
 transcription factors which may control the activity of p53-regulated  
 genes, and are expressed by tumour suppressor genes related to the p53  
 gene family. The gene sequence was isolated from the human colon  
 neuroblastoma cell line IMR-32, using primers V01512-13. The SR-p70d  
 protein sequence is 49 amino acids shorter with a divergence of the  
 first 13 amino acids as compared to the SR-p70a protein (W36184).  
 CC The sequence can be used in the diagnosis and monitoring of cancer,  
 especially neuroblastoma. The nucleic acid sequences and corresponding  
 antisense sequences, are also useful in gene therapy, e.g. to regulate  
 apoptosis.  
 CC Sequence 587 AA;

Query Match 90.4%; Score 3058; DB 1; Length 587;  
 Best Local Similarity 100.0%; Pred. No. 3.3e-259;  
 Matches 574; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

63 AQFNLLSTMDQMSRAASAPYTPHEAASVTPHSPVAQPSSTFTDMSAPVPSNTDYP 122  
 14 AQFNLLSTMDQMSRAASAPYTPHEAASVTPHSPVAQPSSTFTDMSAPVPSNTDYP 73  
 123 GPHFEVTFQOSSTAKSATWTYSPLKKLYCQIAKTCPIQIKVSTPPPGTAIRAMPYK 182  
 74 GPHFEVTFQOSSTAKSATWTYSPLKKLYCQIAKTCPIQIKVSTPPPGTAIRAMPYK 133  
 183 KAEHTVDVVKCPNHELGRDNFEGOSAPASHLIRVEGNLSQYVDDPVTGRQSVVYEP 242  
 134 KAEHTVDVVKCPNHELGRDNFEGOSAPASHLIRVEGNLSQYVDDPVTGRQSVVYEP 193  
 243 POGVTEFTILYFNMCNNSCVGMNRRPILIIITLEMKGQVLRGRSFEGRICACGRDR 302  
 194 POGVTEFTILYFNMCNNSCVGMNRRPILIIITLEMKGQVLRGRSFEGRICACGRDR 253  
 303 KADEHYREQQALNESSAKRAFKQSPAPALGAGVKKRRHGDEDTYLVQYRGR 362  
 254 KADEHYREQQALNESSAKRAFKQSPAPALGAGVKKRRHGDEDTYLVQYRGR 313  
 363 ENFETLMKLSLELMELVQPLVDYSYRQOQQLLQRPSPHLPSPYGPVLSPMKRVHGMN 422  
 314 ENFETLMKLSLELMELVQPLVDYSYRQOQQLLQRPSPHLPSPYGPVLSPMKRVHGMN 373  
 423 KLPVSNQVLGVGPPHSSAATPNLGPVGMNHNHGHAVPANGEMSSSHSAQSMVSGSHCT 482  
 374 KLPVSNQVLGVGPPHSSAATPNLGPVGMNHNHGHAVPANGEMSSSHSAQSMVSGSHCT 433  
 483 PPPPHADPSLVSLTGLGCPNCIEYFTSQGLSIYHLQNTIEDLQALKIPQYRMTIW 542  
 434 PPPPHADPSLVSLTGLGCPNCIEYFTSQGLSIYHLQNTIEDLQALKIPQYRMTIW 493  
 543 RGLQDLKQGHDIYTAQQLLRSSNAATISIGGSGELQQRVNEAVHFRVHTITIPNRRGP 602  
 494 RGLQDLKQGHDIYTAQQLLRSSNAATISIGGSGELQQRVNEAVHFRVHTITIPNRRGP 553  
 603 GGGPDWADFGDLPDCKARKQPIKEEFTAEI 636  
 554 GGGPDWADFGDLPDCKARKQPIKEEFTAEI 587

RESULT 6  
 36185  
 C W36185 standard; Protein; 589 AA.  
 C W36185;  
 T 27-APR-1998 (first entry)

DE Mouse p53 tumour suppressor-related protein SR-p70c.  
 KW SR-p70; mouse; transcription factor; p53; tumour suppressor gene;  
 KW homology; differential splicing; diagnosis; cancer; neuroblastoma;  
 KW gene therapy; apoptosis.  
 PS Mus musculus.  
 PN W09728186-A1.  
 PD 07-AUG-1997.  
 PF 03-FEB-1997; F00214.  
 PR 02-FEB-1996; FR-001309.  
 PA (SNFI) SANOFI SA.  
 PI Caput D, Ferrara P, Kaghad AM;  
 DR WPI; 97-402550/37.  
 DR N-PSDB; V01499.  
 PT New polypeptide(s) encoded by the SR-p70 tumour suppressor gene -  
 and related nucleic acid, useful for diagnosis and treatment of  
 tumours  
 PS Claim 1; Fig 7; 136pp; French.  
 CC This is the amino acid sequence of the mouse protein SR-p70c. SR-p70  
 are transcription factors which may control the activity of p53-regulated  
 genes, and are expressed by tumour suppressor genes related to the p53  
 gene family. The gene sequence was isolated from the mouse pituitary  
 tumour cell line Atr-20, using primers V01508-9. The sequence can be  
 used in the diagnosis and monitoring of cancer, especially neuroblastoma.  
 CC The nucleic acid sequences and corresponding antisense sequences, are  
 also useful in gene therapy, e.g. to regulate apoptosis.  
 CC Sequence 589 AA;

Query Match 83.6%; Score 2830.5; DB 1; Length 589;  
 Best Local Similarity 91.7%; Pred. No. 2.7e-239;  
 Matches 531; Conservative 19; Mismatches 22; Indels 7; Gaps 4;

63 AQFNLLSTMDQMSRAASAPYTPHEAASVTPHSPVAQPSSTFTDMSAPVPSNTDYP 122  
 13 AQFNLLSTMDQMSRAASAPYTPHEAASVTPHSPVAQPSSTFTDMSAPVPSNTDYP 72  
 123 GPHFEVTFQOSSTAKSATWTYSPLKKLYCQIAKTCPIQIKVSTPPPGTAIRAMPYK 182  
 73 GPHFEVTFQOSSTAKSATWTYSPLKKLYCQIAKTCPIQIKVSTPPPGTAIRAMPYK 132  
 183 KAEHTVDVVKCPNHELGRDNFEGOSAPASHLIRVEGNLSQYVDDPVTGRQSVVYEP 242  
 133 KAEHTVDVVKCPNHELGRDNFEGOSAPASHLIRVEGNLSQYVDDPVTGRQSVVYEP 192  
 243 POGVTEFTILYFNMCNNSCVGMNRRPILIIITLEMKGQVLRGRSFEGRICACGRDR 302  
 193 POGVTEFTILYFNMCNNSCVGMNRRPILIIITLEMKGQVLRGRSFEGRICACGRDR 252  
 303 KADEHYREQQALNESSAKRAFKQSPAPALGAGVKKRRHGDEDTYLVQYRGR 362  
 253 KADEHYREQQALNESSAKRAFKQSPAPALGAGVKKRRHGDEDTYLVQYRGR 312  
 363 ENFETLMKLSLELMELVQPLVDYSYRQOQQLLQRPSPHLPSPYGPVLSPMKRVHGMN 420  
 313 ENFETLMKLSLELMELVQPLVDYSYRQOQQLLQRPSPHLPSPYGPVLSPMKRVHGMN 372  
 421 MNKLPVSNQVLGVGPPHSSAATPNLGPVGMNHNHGHAVPANGEMSSSHSAQSMVSGSH 480  
 373 VNKLPVSNQVLGVGPPHSSAAGPNLGPVGMNHNHGHAVPANGEMSSSHSAQSMVSGSH 432  
 481 CTPPPPHADPSLVSLTGLGCPNCIEYFTSQGLSIYHLQNTIEDLQALKIPQYRMT 540  
 433 CTPPPPHADPSLVSLTGLGCPNCIEYFTSQGLSIYHLQNTIEDLQALKIPQYRMT 492  
 541 IWRGLQDLKQGHDIYTAQQLLRSSNAATISIGGSGELQQRVNEAVHFRVHTITIPNR 599  
 493 IWRGLQDLKQGHDIYTAQQLLRSSNAATISIGGSGELQQRVNEAVHFRVHTITIPNR 550  
 600 GGGPDWADFGDLPDCKARKQPIKEEFTAEI 636  
 551 GGGPDWADFGDLPDCKARKQPIKEEFTAEI 589

RESULT 7  
W36190  
ID W36190 standard; Protein; 499 AA.  
AC W36190;  
DC 27-APR-1998 (first entry)  
DE Human p53 tumour suppressor-related protein SR-p70b.  
KW SR-p70; human; transcription factor; p53; tumour suppressor gene;  
KW homology; differential splicing; diagnosis; cancer; neuroblastoma;  
KW gene therapy; apoptosis.  
OS Homo sapiens.  
PN W09728186-A1.  
PD 07-AUG-1997.  
PF 03-FEB-1997; F00214.  
PR 02-FEB-1996; FR-001309.  
PA (SNFI) SANOFI SA.  
PI Caput D, Ferrara P, Kaghad AM;  
DR WPI; 97-402550/37.  
DR N-PSDB; V01505.  
PT New polypeptide(s) encoded by the SR-p70 tumour suppressor gene -  
and related nucleic acid, useful for diagnosis and treatment of  
tumours.  
PS Claim 1; page 72-73; 136pp; French.  
CC This is the amino acid sequence of the human protein SR-p70b. SR-p70 are  
transcription factors which may control the activity of p53-regulated  
genes, and are expressed by tumour suppressor genes related to the p53  
gene family. The gene sequence was isolated from the human neuroblastoma  
cell line SK-N-SH, using primers V01515 and V01518. The SR-p70b gene  
sequence contains a 94 bp deletion between bases 1516-1517 as compared  
to the SR-p70a sequence (V01498). This deletion causes a reading frame  
shift resulting in the generation of a stop codon at position 1498-1500.  
The resultant protein is truncated by 137 amino acids as compared to the  
SR-p70a protein (W36184). The sequence can be used in the diagnosis and  
monitoring of cancer, especially neuroblastoma. The nucleic acid  
sequences and corresponding antisense sequences, are also useful in gene  
therapy, e.g. to regulate apoptosis.  
SQ Sequence 499 AA;

Query Match 77.5%; Score 2624; DB 1; Length 499;  
Best Local Similarity 100.0%; Pred. No. 2.5e-221;  
Matches 494; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 1 MAQSTATSPDGGTTFEHLWSSLEPDSYFDLPQSSRGNNVVGTTDSSMDVHLEGMTTS 60  
DB 1 MAQSTATSPDGGTTFEHLWSSLEPDSYFDLPQSSRGNNVVGTTDSSMDVHLEGMTTS 60  
QY 61 VMAQFNLLSSTMDQSSRAASASPYTPEHAASVTHSPYAQPSSTFTDMSAPVIPSNNTD 120  
DB 61 VMAQFNLLSSTMDQSSRAASASPYTPEHAASVTHSPYAQPSSTFTDMSAPVIPSNNTD 120  
QY 121 YPGPHFEVTFQOSTAKSATWTYSPLLKLYCOIAKTCPIQIKVSTPPPGTAIRAMPY 180  
DB 121 YPGPHFEVTFQOSTAKSATWTYSPLLKLYCOIAKTCPIQIKVSTPPPGTAIRAMPY 180  
QY 181 YKKAHEVTDVVKRCNPHELGRDFNEGOSAPASHLIRVEGNLSQYVDDPVTGROSVVVPY 240  
DB 181 YKKAHEVTDVVKRCNPHELGRDFNEGOSAPASHLIRVEGNLSQYVDDPVTGROSVVVPY 240  
QY 241 EPPQVGTFTILYFNMCNCSVCGMNRRLIITILEMRDQGVLGRRSFEGRICACPGR 300  
DB 241 EPPQVGTFTILYFNMCNCSVCGMNRRLIITILEMRDQGVLGRRSFEGRICACPGR 300  
QY 301 DRKADEHYRQOALNESSAKNGAASKRAFKQSPPAVPAALGAGVKRRHGDDETYIYQVR 360  
DB 301 DRKADEHYRQOALNESSAKNGAASKRAFKQSPPAVPAALGAGVKRRHGDDETYIYQVR 360  
QY 361 GRENFELMKLESLELMELVPQPLVDSYRQOQLLRPSHLQPPSYGVLSPMNKHYGG 420  
DB 361 GRENFELMKLESLELMELVPQPLVDSYRQOQLLRPSHLQPPSYGVLSPMNKHYGG 420  
QY 421 MNKLPSVNLQVQPPHSSAATPNLGPVPGMKNHGHVPAANGEMSSSSHAQSMVSGSH 480  
DB 421 MNKLPSVNLQVQPPHSSAATPNLGPVPGMKNHGHVPAANGEMSSSSHAQSMVSGSH 480

QY 481 CTPPPYPHADPSLV 494  
DB 481 CTPPPYPHADPSLV 494  
RESULT 8  
W36183  
ID W36183 standard; Protein; 499 AA.  
AC W36183;  
DC 27-APR-1998 (first entry)  
DE Monkey p53 tumour suppressor-related protein SR-p70b.  
KW SR-p70; monkey; transcription factor; p53; tumour suppressor gene;  
KW homology; differential splicing; diagnosis; cancer; neuroblastoma;  
KW gene therapy; apoptosis.  
OS Cercopithecus aethiops.  
PN W09728186-A1.  
PD 07-AUG-1997.  
PF 03-FEB-1997; F00214.  
PR 02-FEB-1996; FR-001309.  
PA (SNFI) SANOFI SA.  
PI Caput D, Ferrara P, Kaghad AM;  
DR WPI; 97-402550/37.  
DR N-PSDB; V01497.  
PT New polypeptide(s) encoded by the SR-p70 tumour suppressor gene -  
and related nucleic acid, useful for diagnosis and treatment of  
tumours.  
PS Claim 1; Fig 5; 136pp; French.  
CC This is the amino acid sequence of the protein SR-p70b from monkey cells.  
SR-p70 are transcription factors which may control the activity of  
p53-regulated genes, and are expressed by tumour suppressor genes related  
to the p53 gene family. The gene sequence was isolated from a cDNA  
library by sequencing the inserts and comparing to sequence databases.  
The protein sequence contains regions of homology to the p53 protein.  
The SR-p70b gene sequence was isolated simultaneously with the SR-p70a  
sequence (V01496) from the library and is created by differential  
splicing of the SR-p70 mRNA sequence. The sequences can be used in the  
diagnosis and monitoring of cancer, especially neuroblastoma. The  
nucleic acid sequences and corresponding antisense sequences, are also  
useful in gene therapy, e.g. to regulate apoptosis.  
SQ Sequence 499 AA;

Query Match 76.1%; Score 2574; DB 1; Length 499;  
Best Local Similarity 97.8%; Pred. No. 5.8e-217;  
Matches 483; Conservative 3; Mismatches 8; Indels 0; Gaps 0;  
QY 1 MAQSTATSPDGGTTFEHLWSSLEPDSYFDLPQSSRGNNVVGTTDSSMDVHLEGMTTS 60  
DB 1 MAQSTATSPDGGTTFEHLWSSLEPDSYFDLPQSSRGNNVVGTTDSSMDVHLEGMTTS 60  
QY 61 VMAQFNLLSSTMDQSSRAASASPYTPEHAASVTHSPYAQPSSTFTDMSAPVIPSNNTD 120  
DB 61 VMAQFNLLSSTMDQSSRAASASPYTPEHAASVTHSPYAQPSSTFTDMSAPVIPSNNTD 120  
QY 121 YPGPHFEVTFQOSTAKSATWTYSPLLKLYCOIAKTCPIQIKVSTPPPGTAIRAMPY 180  
DB 121 YPGPHFEVTFQOSTAKSATWTYSPLLKLYCOIAKTCPIQIKVSTPPPGTAIRAMPY 180  
QY 181 YKKAHEVTDVVKRCNPHELGRDFNEGOSAPASHLIRVEGNLSQYVDDPVTGROSVVVPY 240  
DB 181 YKKAHEVTDVVKRCNPHELGRDFNEGOSAPASHLIRVEGNLSQYVDDPVTGROSVVVPY 240  
QY 241 EPPQVGTFTILYFNMCNCSVCGMNRRLIITILEMRDQGVLGRRSFEGRICACPGR 300  
DB 241 EPPQVGTFTILYFNMCNCSVCGMNRRLIITILEMRDQGVLGRRSFEGRICACPGR 300  
QY 301 DRKADEHYRQOALNESSAKNGAASKRAFKQSPPAVPAALGAGVKRRHGDDETYIYQVR 360  
DB 301 DRKADEHYRQOALNESSAKNGAASKRAFKQSPPAVPAALGAGVKRRHGDDETYIYQVR 360  
QY 361 GRENFELMKLESLELMELVPQPLVDSYRQOQLLRPSHLQPPSYGVLSPMNKHYGG 420  
DB 361 GRENFELMKLESLELMELVPQPLVDSYRQOQLLRPSHLQPPSYGVLSPMNKHYGG 420







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1b 228 DCTTHNYMNCSCMGNRRPILITLSSGNLGRNSFEVRCACPGDRRREEE 287
2y 308 HYREQALNESSAKNG-----AASKRAFKQSPAPVAGVKKRRHGHDEDTYYLQV 359
1b 288 NLR-----KKGEPHHELPPGSTRALPNTSSSPQ-----PKKKPLDGEYFTLQI 332
2y 360 RGRENFEILMKLESLELME 379
1b 333 RGRERFEMFRELNEALELKD 352

RESULT 13
ID W13954 standard; Protein; 363 AA.
AC W13954;
DE Modified p53 variant (first entry)
DE Modified p53 variant (del364-393).
KW p53; tumour suppressor; cancer; therapy; cell proliferation;
KW apoptosis; protein engineering; DNA binding.
CS Synthetic.
PN W09710843-A1.
PF 20-SEP-1996; U15188.
PR 22-SEP-1995; US-004802.
PR 21-AUG-1996; US-697221.
PR (WIST-) WISTAR INST ANATOMY & BIOLOGY.
PR Halazometis TD;
PR WPI; 97-202618/18.
PT R284K modified p53 protein having DNA binding ability - useful in
PT treatment of cancer
PT Example 1; 49-51; 82pp; English.
CC A modified p53 variant (W13954) comprises wild-type p53 (see
CC also W13948) having a deletion of the C-terminal 30 amino acids,
CC and is obt'd. by site-directed mutagenesis of p53 DNA. Deletion of
CC the p53 C-terminal 30 amino acids activates the DNA binding of
CC common Class I p53 mutants (see also W13951-52). Novel modified
CC p53 variants (W13949-50, W13953-54, W13968-77), some contg.
CC C-terminal deletions, provide the means for pharmacological rescue
CC of p53 function in cancer patients. Nucleic acids coding for
CC modified p53 can be used for cancer gene therapy.
CC Sequence 363 AA;

Query Match 21.4%; Score 724.5; DB 1; Length 363;
Best Local Similarity 42.1%; Pred. No. 2.5e-55;
Matches 160; Conservative 58; Mismatches 103; Indels 59; Gaps 10;

2y 14 TFEHLWSSLEPSTVFDLPQSSRGNNVGGTDSMDVHFLEGMTTSVMAQFNLLSMD 73
1b 18 TFSDLWKLLENVLSPLP-----SQAMDDLMLSPDD-----IEQWTFDPPGD 61
2y 74 QMSRAASAPVTPHEAASVPTSHSYPAPSSFTDMSAP-----VIPSNTDYPGPHF 127
1b 62 EAPRMPFAAPPVAPAPATP-----AAP-----APAPSWPLSSVPSQKTVGSVGF 109
2y 128 EYTFQOSSTAKSATWTYSPLLKLYCQIAKTCPIQIKVSTPPPGTAIRAMPVYKAEHV 187
1b 110 RLGLFHSHTAKSVTCYSPALNKMFCQLAKTCVQLWVDSSTPPGTRVRAMAIYKQSHM 169
2y 188 TDVVKCPNHELGRDNFNEGQSPASHLIRVEGNNSQYVDDPVTGROSVVYVPEPQVGT 247
1b 170 TEVVRCPHHERGSD--SDG--LAPPQHLIRVEGNLRVEYLDLDRNTFRHSVVVPEPEVGS 227
2y 248 EFTTILYFMNCSSCVGGMNRRPILITILEMRDGOVLGRRRFEGRICACPGDRKAD 307
1b 228 DCTTHNYMNCSCMGNRRPILITLSSGNLGRNSFEVRCACPGDRRREEE 287
2y 308 HYREQALNESSAKNG-----AASKRAFKQSPAPVAGVKKRRHGHDEDTYYLQV 359
1b 288 NLR-----KKGEPHHELPPGSTRALPNTSSSPQ-----PKKKPLDGEYFTLQI 332
2y 360 RGRENFEILMKLESLELME 379

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Db 333 RGRERFEMFRELNEALELKD 352

RESULT 14
ID R22238 standard; Protein; 393 AA.
AC R22238;
DE 23-JUL-1992 (first entry)
DE Sequence of 53 kD cellular protein.
KW Cancer therapy; cancer suppressor gene; oncogenesis.
OS Homo sapiens.
PN EP-475623-A.
PD 18-MAR-1992.
PF 23-AUG-1991; 307791.
PR 24-AUG-1990; US-573405.
PR (REGC) UNIV OF CALIFORNIA.
PI Lee WH, Chen PL;
DR WPI; 92-090221/12.
DR N-PSDB; Q22995.
PT Cloned p53 cDNA and protein prods. - for suppression of
PT neoplastic phenotype e.g. in osteo-sarcoma(s), leukaemia(s),
PT lymphoma(s), etc. English.
PS Claim 2; Page 14; 25pp; English.
CC p53 cDNA, or its gene prods., can be used to suppress and eradicate
CC cancers caused by defective, mutant or absent cancer suppressor
CC genes. Variant forms of p53 are found in human breast, lung or
CC colon carcinoma, lymphoma, leukaemia, etc., suggesting that mutation
CC of the p53 genes is involved in oncogenesis. Specifically 273 Arg
CC is replaced by 273 His, a mutation found exclusively in tumour cells.
CC Sequence 393 AA;

Query Match 21.4%; Score 724.5; DB 1; Length 393;
Best Local Similarity 41.0%; Pred. No. 2.8e-55;
Matches 157; Conservative 57; Mismatches 104; Indels 65; Gaps 9;

QY 14 TFEHLWSSLEPSTVFDLPQSSRGNNVGGTDSMDVHFLEGMTTSVMAQFNLLSMD 73
1b 18 TFSDLWKLLENVLSPLP-----SQAMDDLMLSPDDIE 51
QY 74 QMSRAASAPVTPHEAASVPTSHSYPAPSSFTDMSAPV-----IPSNTRYDGP 124
1b 52 QMFTEDPG-----PDEAPRMPFAAPPVAPAPATP-----APAPSWPLSSVPSQKTVGS 106
QY 125 HFEVTFQOSSTAKSATWTYSPLLKLYCQIAKTCPIQIKVSTPPPGTAIRAMPVYKKA 184
1b 107 YGRLGFLHSHTAKSVTCYSPALNKMFCQLAKTCVQLWVDSSTPPGTRVRAMAIYKQS 166
QY 185 EHVTDVVKCPNHELGRDNFNEGQSPASHLIRVEGNNSQYVDDPVTGROSVVYVPEPQ 244
1b 167 QHMTVEVVRCPHHERGSD--SDG--LAPPQHLIRVEGNLRVEYLDLDRNTFRHSVVVPEPE 224
QY 245 VGTFTTILYFMNCSSCVGGMNRRPILITILEMRDGOVLGRRRFEGRICACPGDRKA 304
1b 225 VGSDCITTHNYMNCSCMGNRRPILITLSSGNLGRNSFEVRCACPGDRRT 284
QY 305 DEDHYEQALNESSAKNG-----AASKRAFKQSPAPVAGVKKRRHGHDEDTYY 356
1b 285 EENLR-----KKGEPHHELPPGSTRALPNTSSSPQ-----PKKKPLDGEYFT 329
QY 357 LQVRGENFEILMKLESLELME 379
1b 330 LQIRGRERFEMFRELNEALELKD 352

RESULT 15
ID R26758 standard; peptide; 393 AA.
AC R26758;
DE 09-FEB-1993 (first entry)
DE p53.
KW Point mutation; translocation; proto-oncogene; cancer;

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OS antigen-presenting cell; T-cell; HLA.  
FH Synthetic.  
Key Location/Qualifiers  
FT misc\_difference 273  
FT /label= mutation  
FT /note= "Arg -> any amino acid except Arg, see CC"  
PN GB2253211-A.  
PD 02-SEP-1992.  
PD 26-FEB-1992; 004098.  
PR 26-FEB-1991; GB-003974.  
PA (NHSD ) NORSK HYDRO AS.  
PI Eriksen JA, Gaudernack G, Gedde Dahl T;  
DR WPI: 92-294575/36.  
PT Synthetic oncogene-protein peptide - used for treating and  
PT vaccinating against cancers  
PS Disclosure; Page 10; 78pp; English.  
CC New peptides, which have a point mutation or translocation compared to  
CC the corresp. fragment of the proto-oncogene prod., correspond to,  
CC completely cover or are active fragments of a processed oncogene protein  
CC fragment as presented by a cancer cell or other antigen-presenting cell  
CC and are capable of inducing a specific T-cell response to the actual  
CC oncogene protein fragment as produced by the cell and processed and  
CC presented in the HLA mol.  
CC For example, a peptide fragment of p53 comprising at least  
CC mutations in position 273, in which position any amino acid except Arg  
CC may be located. The p53 sequence below was not disclosed in the  
CC specification, but retrieved by the Indexer from Swiss-prot P04637.  
SQ Sequence 393 AA;  
  
Query Match 21.4%; Score 724.5; DB 1; Length 393;  
Best Local Similarity 42.1%; Pred No. 2.8e-55;  
Matches 160; Conservative 58; Mismatches 103; Indels 59; Gaps 10;  
  
QY 14 TFEHLMSSLEPSTYFDLPQSRGNEVYGGTSSMDVPHLEGMTTSVMAQFNLLSSTMD 73  
DB 14 TFEHLMSSLEPSTYFDLPQSRGNEVYGGTSSMDVPHLEGMTTSVMAQFNLLSSTMD 73  
QY 18 TTSDLKWLKLPENNVLSPLP--SQAMDDLMLSPDD-----IEQWFTEDPGPD 61  
DB 18 TTSDLKWLKLPENNVLSPLP--SQAMDDLMLSPDD-----IEQWFTEDPGPD 61  
QY 74 QMSRRAASPTPEHAASVPHSPYAPQSSFTDMSAP-----VIPSNTDYPGPHF 127  
DB 74 QMSRRAASPTPEHAASVPHSPYAPQSSFTDMSAP-----VIPSNTDYPGPHF 127  
QY 62 EAPRMEAPAPPVAPAPAAATP-----AAP-----APAPSWPLSSVPSQTYOGSYGF 109  
DB 62 EAPRMEAPAPPVAPAPAAATP-----AAP-----APAPSWPLSSVPSQTYOGSYGF 109  
QY 128 EYTFQOSTAKSATWTYSPLLKLYCQIAKTCPIQKVSTPPPPGTATRAAMPVYKKAHV 187  
DB 128 EYTFQOSTAKSATWTYSPLLKLYCQIAKTCPIQKVSTPPPPGTATRAAMPVYKKAHV 187  
QY 110 RLGFHSGTAKSVTCYSPALNKMFCQLAKTCFVLWVDSTPPGTRVRAAIYKQSQHM 169  
DB 110 RLGFHSGTAKSVTCYSPALNKMFCQLAKTCFVLWVDSTPPGTRVRAAIYKQSQHM 169  
QY 188 TDVVKPCPNHELGRDNEGOSAPASHLIRVEGNLSQYVDDPVTGRQSVVYVPEPQVGT 247  
DB 188 TDVVKPCPNHELGRDNEGOSAPASHLIRVEGNLSQYVDDPVTGRQSVVYVPEPQVGT 247  
QY 170 TEVVRCPHERCSD-SDG-LAPPQHLIRVEGNLRYEYLDLDRNTFRHSVYVPEPEVGS 227  
DB 170 TEVVRCPHERCSD-SDG-LAPPQHLIRVEGNLRYEYLDLDRNTFRHSVYVPEPEVGS 227  
QY 248 EETILYNFNCNCCVCGMNRPIIITILEMRDGOVLGRSFEGRICACGRDRKADED 307  
DB 248 EETILYNFNCNCCVCGMNRPIIITILEMRDGOVLGRSFEGRICACGRDRKADED 307  
QY 228 DCTIHYNYCNSCGMNRPIIITILEMRDGOVLGRSFEGRICACGRDRKADED 287  
DB 228 DCTIHYNYCNSCGMNRPIIITILEMRDGOVLGRSFEGRICACGRDRKADED 287  
QY 308 HYREQAALNESSAKNG-----AASKRAFKQSPAPVAPALGAGVKKRHGDEDTYILQV 359  
DB 308 HYREQAALNESSAKNG-----AASKRAFKQSPAPVAPALGAGVKKRHGDEDTYILQV 359  
QY 288 NLR-----KCEPHHELPPGSKRALPNTSSSPQ-----PKKPLDGEYFTLQI 332  
DB 288 NLR-----KCEPHHELPPGSKRALPNTSSSPQ-----PKKPLDGEYFTLQI 332  
QY 360 RGRNFELMKLESLELME 379  
DB 360 RGRNFELMKLESLELME 379  
QY 333 RGRFERFRELNEALELKD 352  
DB 333 RGRFERFRELNEALELKD 352